

two substituents R^{1-2} , together with the carbon atoms to which they are attached, form a cycloalkyl or heterocyclcyl which can be substituted by 0, 1 or 2 substituents R^{1-2-1} , the substituents R^{1-2-1} being selected independently of one another from the group consisting of halogen, nitro, amino, trifluoromethyl, hydroxyl, alkyl and alkoxy,

R^2 is hydrogen or methyl,

R^3 is hydrogen, hydroxyl, amino, C_1-C_3 alkyl, benzyl, C_1-C_3 alkoxy, benzyloxy, C_1-C_3 alkylamino, C_1-C_3 alkylcarbonylamino, phenylcarbonylamino or benzylcarbonylamino,

R^4 is hydrogen or C_1-C_3 alkyl,

R^5 is halogen, trifluoromethyl, trifluoromethoxy, nitro, amino, alkylamino, hydroxyl, alkyl, alkoxy, carboxyl, alkoxycarbonyl, aminocarbonyl, alkylaminocarbonyl, aryl or heteroaryl,

or

two substituents R^5 together with the carbon atoms to which they are attached form a cycloalkyl or heterocyclcyl each of which may be substituted by 0, 1 or 2 substituents R^{5-1} , the substituents R^{5-1} being selected independently of one another from the group consisting of halogen, nitro, amino, trifluoromethyl, hydroxyl, alkyl and alkoxy,

R^6 is alkyl, cycloalkyl, cycloalkenyl or heterocyclcyl,

it being possible for R^6 to be substituted by 0, 1 or 2 substituents R^{6-1} , the substituents R^{6-1} being selected independently of one another from the group consisting of halogen, nitro, amino, trifluoromethyl, hydroxyl, alkyl and alkoxy,

n is a number 0, 1, 2 or 3,

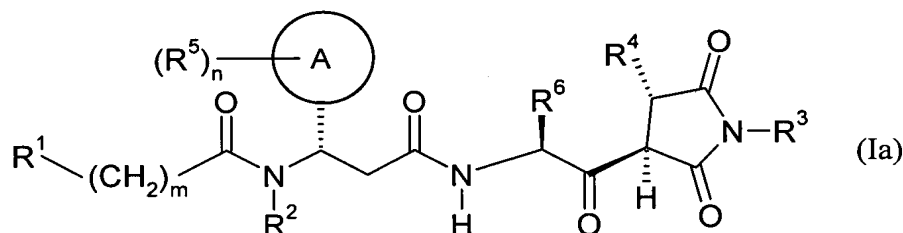
it being possible for the radicals R^5 to be identical or different when n is 2 or 3,

m is a number 0, 1, 2, 3 or 4,

A is aryl or heteroaryl,

or a salt thereof, a solvate thereof or a solvate of a salt thereof.

2. (Previously Presented) A compound according to claim 1, characterized in that it corresponds to the formula



in which R^1 to R^6 , A , m and n have the same definition as in formula (I).

3. (Previously Presented) A compound according to claim 1, characterized in that

R^1 is pyridyl, imidazolyl, thienyl, furyl, oxadiazolyl, pyrazolyl, pyrazinyl, pyridazinyl, pyrimidinyl, quinolinyl or isoquinolinyl,

where R^1 can be substituted by 0, 1 or 2 substituents R^{1-1} , the substituents R^{1-1} being selected independently of one another from the group consisting of halogen, alkyl, amino, trifluoromethyl, phenyl and alkoxy,

or

R^1 is phenyl or naphthyl,

where phenyl or naphthyl are substituted by 1, 2 or 3 substituents R^{1-2} , the substituents R^{1-2} being selected independently of one another from the group consisting of halogen, C_1 - C_4 alkyl, dimethylamino, cyano, trifluoromethyl, 3- to 7-membered cycloalkyl, 5- or 6-membered heterocyclyl, phenyl, 5- or 6-membered heteroaryl, C_1 - C_3 alkoxy, phenoxy, benzyloxy, phenylcarbonylamino and aminosulfonyl,

or

two substituents R^{1-2} , together with the carbon atoms to which they are attached, form a 1,3-benzodioxole or a 1,4-benzodioxane,

R^2 is hydrogen,

R^3 is hydrogen, amino, methyl, methoxy, ethoxy, methylamino or dimethylamino,

R^4 is methyl,

R^5 is fluoro, chloro, trifluoromethyl, C_1 - C_4 alkoxy, methoxycarbonyl, C_1 - C_4 alkyl, phenyl or pyridyl,

or

two substituents R^5 , together with the phenyl ring to which they are attached, form a 1,3-benzodioxole or a 1,4-benzodioxane,

R^6 is C_3 - C_6 alkyl or 3- to 6-membered cycloalkyl,

n is a number 0, 1 or 2,

and, if n is 2, the radicals R^5 can be identical or different,

m is a number 0, 1, 2 or 3,

and

A is phenyl, naphthyl, pyridyl, thienyl, furanyl, quinolinyl or isoquinolinyl.

4. (Previously Presented) A compound according to claim 1, characterized in that

R^1 is pyridyl, thienyl, furyl, quinolinyl or isoquinolinyl,

where R^1 can be substituted by 0, 1 or 2 substituents R^{1-1} , the substituents R^{1-1} being selected independently of one another from the group consisting of halogen, C_1 - C_4 alkyl, trifluoromethyl, phenyl and C_1 - C_3 -alkoxy,

or

R^1 is phenyl or naphthyl,

where phenyl or naphthyl are substituted by 1, 2 or 3 substituents R^{1-2} , the substituents R^{1-2} being selected independently of one another

from the group consisting of halogen, C₁-C₄ alkyl, dimethylamino, cyano, trifluoromethyl, 5- or 6-membered heterocyclyl, 5- or 6-membered heteroaryl, C₁-C₃ alkoxy, phenyloxy or benzyloxy,

or

two substituents R¹⁻², together with the carbon atoms to which they are attached, form a 1,3-benzodioxole or a 1,4-benzodioxane,

R² is hydrogen,

R³ is hydrogen, amino, methylamino or dimethylamino,

R⁴ is methyl,

R⁵ is fluoro, chloro, trifluoromethyl, C₁-C₃ alkoxy, C₁-C₄ alkyl, phenyl or pyridyl,

R⁶ is isopropyl, tert-butyl, isopentyl, cyclopentyl or cyclohexyl,

n is a number 0, 1 or 2,

and, if n is 2, the radicals R⁵ can be identical or different,

m is a number 0, 1 or 2,

and

A is phenyl, naphthyl, pyridyl, thienyl, quinolinylnyl or isoquinolinylnyl.

5. (Previously Presented) A compound according to claim 1, characterized in that

R¹ is pyridyl, thienyl, furyl, quinolinylnyl or isoquinolinylnyl,

where R¹ can be substituted by 0, 1 or 2 substituents R¹⁻¹, the substituents R¹⁻¹ being selected independently of one another from the group consisting of fluoro, chloro, trifluoromethyl, C₁-C₄ alkyl, phenyl and methoxy.

6. (Previously Presented) A compound according to claim 1, characterized in that

R^1 is phenyl or naphthyl,

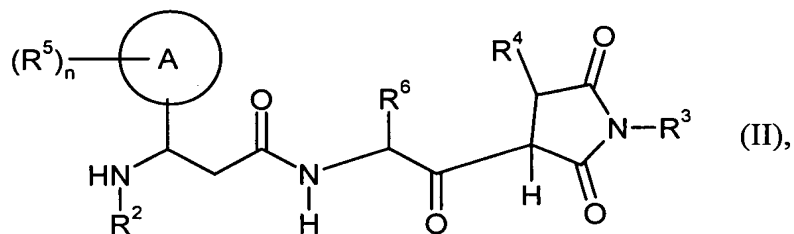
where phenyl or naphthyl are substituted by 1, 2 or 3 substituents R^{1-2} , the substituents R^{1-2} being selected independently of one another from the group consisting of halogen, C_1 - C_4 alkyl, dimethylamino, cyano, trifluoromethyl, 5- or 6-membered heterocyclyl, 5- or 6-membered heteroaryl, C_1 - C_3 alkoxy, phenyloxy or benzyloxy,

or

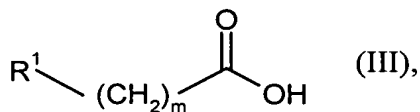
two substituents R^{1-2} , together with the carbon atoms to which they are attached, form a 1,3-benzodioxole or a 1,4-benzodioxane.

7. (Previously Presented) A compound according to claim 1, characterized in that R^2 is hydrogen.
8. (Previously Presented) A compound according to claim 1, characterized in that R^3 is hydrogen or amino.
9. (Previously Presented) A compound according to claim 1, characterized in that R^4 is methyl.
10. (Previously Presented) A compound according to claim 1, characterized in that n is the number zero.
11. (Previously Presented) A compound according to claim 1, characterized in that n is the number 1, A is phenyl and R^5 is fluoro, chloro, trifluoromethyl, alkoxy, C_1 - C_4 -alkyl, phenyl or pyridyl, R^5 being positioned meta or para to the linkage site of the phenyl ring.
12. (Previously Presented) A compound according to claim 1, characterized in that R^6 is C_3 - C_6 -alkyl or 3- to 6-membered cycloalkyl.
13. (Previously Presented) A compound according to claim 1, characterized in that m is the number zero.
14. (Previously Presented) A compound according to claim 1, characterized in that A is phenyl, naphthyl, pyridyl, thienyl, quinolinyl or isoquinolinyl.
15. (Previously Presented) A process for preparing a compound of formula (I) according to claim 1, characterized in that

[A] a compound of the formula



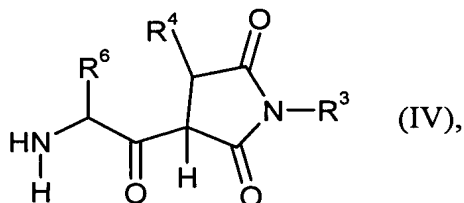
in which R^2 to R^6 , A and n are as defined in claim 1, is reacted with a compound of the formula



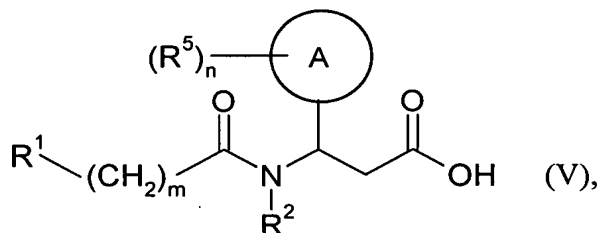
in which R^1 and m are as defined in claim 1,

or

[B] a compound of the formula



in which R^3 , R^4 and R^6 are as defined in claim 1, is reacted with a compound of the formula



in which R^1 , R^2 , R^5 , A, m and n are as defined in claim 1.

16. (Canceled)

17. (Currently Amended) A ~~medicinal product~~ pharmaceutical composition comprising at least one compound of claim 1 in combination with at least one ~~pharmaceutically compatible~~, pharmaceutically acceptable carrier or ~~other excipients~~ excipient.
18. (Canceled)
19. (Canceled)
20. (Currently Amended) A method of controlling bacterial infections in ~~people and animals by a person or an animal comprising administering to a person or animal, in need thereof,~~ an antibacterially effective amount of at least one compound of claim 1.
21. (New) A method of controlling bacterial infections in a person or an animal comprising administering to a person or animal, in need thereof, an antibacterially effective amount of at least one composition of claim 17.